07 C 6561

JUDGE NORDBERG MAGISTRATE JUDGE SCHENKIER

GROUP EXHIBIT A



Notice of Service of Process

MIW / ALL

Transmittal Number: 5405360 **Date Processed: 10/23/2007**

Primary Contact:

Jeffery Kruse Boston Scientific/Guidant 4100 Hamline Avenue North

St. Paul, MN 55112

Copy of transmittal only provided to:

Brenda McKee Diane Barker Julie Somora Kelly Phillips Mildred Helgeson

Entity:

Guidant Corporation

Entity ID Number 2488008

Entity Served:

Guidant Corporation

Title of Action:

Judith Maher vs. Guidant Corporation

Document(s) Type:

Summons/Complaint

Nature of Action:

Product Liability

Court:

Cook Circuit Court, Illinois

Case Number:

2007 L007363

Jurisdiction Served:

Oregon

Date Served on CSC:

10/22/2007

Answer or Appearance Due:

30 Days

Originally Served On:

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How Served:

Regular Mail

Plaintiff's Attorney:

Scott A. Kogen 312-782-7341

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THE LAW OFFICES OF SCOTT A. KOGEN & ASSOCIATES, PC.

Attorneys at Law
134 N. LaSalle Street, Suite 1515
Chicago, Illinois 60602
(312) 782-7341
(312) 377-1771 Facsimile

October 18, 2007

VIA REGULAR MAIL

Guidant Corporation 285 Liberty Street Salem, Nevada 97301 Attn: Claims Department

Re: Judith Maher v. Guidant Corporation et. al.

Case Number: 2007 L 007363

Dear Claims Department,

Enclosed is a copy of the Complaint and summons issued in the case against Guidant. Please send me a copy of your appearance in this matter as soon as possible

Sincerely,

Scott A. Kogen

SAK/dmj

Encs.

2220 - Not Served 2320 - Served By Mail 2420 - Served By Publication SUMMONS	2221 - Served 2221 - Not Served 2321 - Served By Mail 2421 - Served By Publication ALIAS - SUMMONS	CCG N001-10M-1	I-07-05 (
	THE CIRCUIT COURT OF CO	OOK COUNTY, ILL	INOIS DIVISION
(Name all parties)			
Judith Maker		_	;
Guidant Corporation C	v. and Guident Sales Corp	No. <u>2</u> See	2007 L 007362 Service List
	SUMMO	NS	
To each Defendant:			
following location:	e your appearance, and pay the r	equired fee, in the Of	t in this case, a copy of which is fice of the Clerk of this Court at the
Richard J. Daley Ce	nter, 50 W. Washington, Room	<u>802</u> , Ch	icago, Illinois 60602
District 2 - Skokie 5600 Old Orchard Re Skokie, IL 60077	District 3 - Rolling I 2121 Euclid Rolling Meadows, I	_	District 4 - Maywood 1500 Maybrook Ave. Maywood, IL 60153
District 5 - Bridgevie 10220 S. 76th Ave. Bridgeview, IL 60455	16501 S. Kedzie Pky	vy.	Child Support 28 North Clark St., Room 200 Chicago, Illinois 60602
You must file within 30 days after IF YOU FAIL TO DO SO, A JUREQUESTED IN THE COMP	UDUMENT RV HERALIT MA	ounting the day of an	
To the officer:			
This Summons must be endorsement of service and feet be returned so endorsed. This St			m it was given for service, with not be made, this Summons shall date.
Atty. No.: 32193		WITNESS	OCT 1,8 2007
Name: Lin Offices of Soft Atty. for: Phintelf	1 kgen & Asocites, 80.		THE TAX BROWN
Address: 134 N. Visalle Street	it so he kik		Clerk of Court
City/State/Zip: Chrack, Il	1612	Date of service:	• • • • • • • • • • • • • • • • • • • •
Telephone: <u>(312) 7.62-7.34/</u>	/	(To be inserted by or oth	officer on copy left with defendant ner person)
Service by Facsimile Transmission	on will be accepted at:	(Area Coda) (Francis	A A

DOROTHY BROWN, CLERK OF THE CIRCUIT COURT OF COOK COUNTY, ILLINOIS

(Area Code) (Facsimile Telephone Number)

IN THE CIRCUIT COURT OF COOK COUNTY, ILLINOIS COUNTY DEPARTMENT, LAW DIVISION

JUDITH MAHER,)	
Plaintiff,)	
v.)	No.
GUIDANT CORPORATION, and GUIDANT SALES CORPORATION,)	
Defendants.)	

NOW COMES Plaintiff, JUDITH MAHER, by and through his attorneys, LAW OFFICES OF SCOTT A. KOGEN & ASSOCIATES, P.C., and complains of the Defendants, as follows:

COMPLAINT AT LAW

PARTIES

- 1. Defendant Guidant Corporation is a publicly traded corporation and has it principal place of business in Indianapolis, Indiana. Guidant Corporation designs, manufactures, and markets the product that is at issue in this Complaint.
- 2. Defendant Guidant Sales Corporation is a wholly owned subsidiary of Defendant Guidant Corporation and has its principal place of business in Indianapolis, Indiana. Guidant Sales Corporation markets, distributes, and sells the product that is at issue in this Complaint.
- Plaintiff, Judith Maher lives in Chicago, IL, and had a Guidant Discovery, Model
 serial number 403115, implanted in September of 1998.

GENERAL ALLEGATIONS

- 4. Plaintiff realleges Paragraphs 1 through 3 as though fully stated herein.
- 5. Defendants designed, tested, manufactured, and provided labeling for implantable pacemakers, including Discovery, Model 1274.
- 6. These devices are surgically implanted in persons for certain types of heart disease that creates the risk of a life-threatening heart arrhythmia.
- 7. Plaintiff had Discovery, Model 1274, Serial Number 403115 implanted into her chest. Due to information regarding the defective nature of this pacemaker, Plaintiff had it removed on October 19, 2005.
- 8. The Discovery, Model 1274, manufactured between November 25, 1997 and October 26, 2000, was recalled on July 18, 2005, by Defendants. The seal within the devices can leak, allowing moisture to affect the electronic circuits. This defect can cause the pacemakers to fail to provide pacing or can cause a rapid heart rate or other unexpected side effects. The problems may occur without warning and can lead to loss of consciousness, and possibly heart failure and death.
- 9. The device does not give any sign of impending failure. There is no test that could have predicted whether Plaintiff, Judith Maher's device would fail.
- 10. Plaintiff did not discover the defect until July 22, 2005, when she received her recall letter from Defendants.

COUNT I- NEGLIGENCE

11. Plaintiff realleges Paragraphs 1 through 10 as though fully alleged herein.

- 12. Defendants carelessly designed, tested, manufactured, marketed, distributed and sold the DISCOVERY. Model 1274.
- 13. Defendants were negligent in manufacturing the Discovery pacemaker
- 14. Defendants, GUIDANT CORPORATION and GUIDANT SALES
 CORPORATION, breached their duty to Plaintiff through one or more of the following acts and/or omissions:
 - a) Negligently designed, inspected, tested and manufactured the equipment it sold;
 - b) Negligently sold, leased or distributed a defective implantable pacemaker;
 - c) Failed to ensure that said pacemakers were functioning properly prior to sale or distribution.
 - d) Failed to properly test its implantable pacemakers to ensure their safe condition prior to sale or distribution.
- 15. Defendants admitted that the Discovery Model 1274 was dangerous and defective when it recalled this product. Defendants had this actual knowledge on or before July 22, 2005.
- 16. That as a direct and proximate result of one or more of the aforesaid negligent acts/or omissions, Plaintiff, JUDITH MAHER, sustained severe and permanent bodily injury. Further, Plaintiff has suffered great pain and anguish, of both mind and body, and will, in the future, continue to suffer. Plaintiff expended and became liable for and will expend and become liable for large sums of money for medical care and services.

WHEREFORE, Plaintiff, JUDITH MAHER, respectfully requests that this Court enter Judgment in her favor in an amount in excess of \$50,000.00 and costs of this suit.

COUNT II- STRICT LIABILITY

- 17. Plaintiff realleges Paragraphs 1 through 16 as though fully alleged herein
- 18. Defendants, designed, tested, manufactured, marketed, distributed and sold the Discovery pacemaker, Model 1274 in a condition which rendered it unreasonably dangerous due to its propensity to fail without warning.
- 19. At all relevant times material to this matter, Defendants, had a duty to design, market, test, manufacture and/or produce equipment that was safe to individuals using the product for its intended purpose, and refrain from putting in the stream of commerce a defectively manufactured product which was likely to cause serious injury or death to individuals using such product for its intended purpose.
- 20. Defendants, breached their duty to Plaintiff through one or more of the following acts and/or omissions:
 - a) Designed an implantable pacemaker that was defective in that the seal could leak allowing moisture to affect the electrical circuits.
- b) Designed an implantable pace maker that lacked a means of warning the individual in the event that it malfunctioned;
 - c) Designed an implantable pacemaker that was otherwise unsafe and defective;

- Manufactured an implantable pacemaker that lacked adequate safeguards d) and safety devices to prevent malfunctions;
 - e) Failed to properly test the design of the implantable pacemaker prior to placing it into the stream of commerce
 - f) Placed into the stream of commerce an implantable pacemaker that lacked adequate safeguards and safety devices to prevent malfunctions;
 - Placed into the stream of commerce an implantable pacemaker that was g) otherwise unsafe and defective.
 - h) Was otherwise negligent in the manufacturing, design and distribution of said implantable pacemaker.
- 21. That as a direct and proximate result of one or more of the aforesaid product defects, Plaintiff, JUDITH MAHER, suffered great pain and anguish, of both mind and body, and will, in the future, continue to suffer. Plaintiff expended and became liable for and will expend and become liable for large sums of money for medical care and services.
- 22. Defendants admitted that the Discovery Model 1274 was dangerous and defective when it recalled this product. Defendants had this actual knowledge on or before July 22, 2005.
- 23. That the defects existed when Defendants placed these devices into the stream of commerce.
- 24. That as a direct and proximate result of one or more of the aforesaid negligent acts/or omissions, Plaintiff, JUDITH MAHER, sustained severe and permanent bodily

injury. Further, Plaintiff has suffered great pain and anguish, of both mind and body, and will, in the future, continue to suffer. Plaintiff expended and became liable for and will expend and become liable for large sums of money for medical care and services.

WHEREFORE, Plaintiff, JUDITH MAHER, demands judgment against the Defendants, in a sum greater than \$50,000.00 plus costs and any additional relief this Court deems fair and just.

COUNT III BREACH OF WARRANTY

- 25. Plaintiff realleges 1 Paragraphs 24 through as though fully alleged herein.
- 26. Defendants impliedly warranted that its implantable pacemaker devices were merchantable, fit and safe for ordinary use and for a particular purpose.
- 27. Defendants breached all warranties by selling their implantable pacemakers which were defective and unsafe.
- 28. That as a direct and proximate result of one or more of the aforesaid negligent acts/or omissions, Plaintiff, JUDITH MAHER, sustained severe and permanent bodily injury. Further, Plaintiff has suffered great pain and anguish, of both mind and body, and will, in the future, continue to suffer. Plaintiff expended and became liable for and will expend and become liable for large sums of money for medical care and services.

WHEREFORE Plaintiff, JUDITH MAHER, respectfully requests that this Honorable Court enter a Judgment in her favor in an amount exceeding \$50,000.00 and the costs of this lawsuit.

Scott A. Kogen, P.C.

Law Offices of Scott A. Kogen & Associates, P.C. 134 N. LaSalle St., Suite 1515 Chicago, IL 60602 312-782-7341 32193

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Scott A. Kogen & Associates P.C 134 N. LaSalle Suite 1515 Chicago, IL 60502

Guidant Corporation 285 Liberty Street Salem, Nevada 97301

Attn: Claims Department

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